

AMENDMENTS TO THE CLAIMS:

This listing of claims will replace all prior versions, and listings, of claims in the application:

Listing of Claims:

1. (Currently amended): A protein chip reagent utilizing a cell-free protein synthesis system which comprises the following elements:

a: a translation reaction solution containing cell extract for a cell-free protein synthesis comprising a wheat embryo extract wherefrom endosperm components and low-molecular weight protein synthesis inhibitors are substantially removed, substances necessary for protein synthesis containing a substrate and an energy source, and a specific translation template is added for adding to each one or more different well wells of a container which is partitioned in plural sections;

b: ~~at least the substances necessary for protein synthesis containing substrate and energy source and a specific translation template are added to the well mentioned in "a"; and wherein~~

e: the protein chip reagent solution in the well mentioned in "b" is a freeze-dried preparation, prepared by freeze drying (~~by freeze drying~~).

2. (Currently amended): A protein chip reagent utilizing a cell-free protein synthesis system which comprises the following elements:

a: a translation reaction solution containing cell extract for a cell-free protein synthesis comprising a wheat embryo extract wherefrom endosperm components and low-molecular weight protein synthesis inhibitors are substantially removed, substances necessary for protein synthesis containing substrate and energy source, a specific translation template, a protein, and a deliquescent substance is added for adding to each one or more different well wells of a container

which is partitioned in plural sections;

~~b: at least the substances necessary for protein synthesis containing substrate and energy source and a specific translation template are added to the well mentioned in "a"; wherein~~

~~e: the solution in the well mentioned in "b" the protein chip reagent is a freeze-dried preparation, prepared by freeze drying (by freeze drying); and~~

~~d: an amount of a the deliquescent substance in the freeze-dried preparation in the well mentioned in "c" is less than 0.01 part by weight or less to 1 part by weight of the protein in the said freeze-dried preparation.~~

3. (Currently amended): A protein chip reagent utilizing a cell-free protein synthesis system which comprises the following elements:

a: a translation reaction solution containing cell extract for a cell-free protein synthesis comprising a wheat embryo extract wherefrom endosperm components and low-molecular weight protein synthesis inhibitors are substantially removed, substances necessary for protein synthesis containing a substrate and an energy source, a specific translation template, a protein, and a deliquescent substance, is for added adding to each different well wells of a container which is partitioned in plural sections;

~~b: at least the substances necessary for protein synthesis containing substrate and energy source and a specific translation template are added to the well mentioned in "a"; wherein~~

~~e: the solution in the well mentioned in "b" protein chip reagent is a freeze-dried preparation prepared by freeze drying (by freeze drying);~~

~~d: an amount of a the deliquescent substance in the freeze-dried preparation in the well mentioned in "c" is less than 0.01 part by weight or less to 1 part by weight of the protein in the said freeze-dried preparation; and~~

~~e: different kind type of translation template is a plurality of different translation templates are contained in each of the different wells of the container the solution mentioned in "b" and makes for making two or more kinds of proteins synthesizable in each different well of the container which is partitioned in plural sections.~~

4. (Currently amended): A protein chip reagent utilizing a cell-free protein synthesis system which comprises the following elements:

a: a translation reaction solution containing cell extract for a cell-free protein synthesis comprising a wheat embryo extract wherefrom endosperm components and low-molecular weight protein synthesis inhibitors are substantially removed, substances necessary for protein synthesis containing substrate and energy source, a specific translation template, a protein, and a deliquescent substance is added to each for adding to one or more different well wells of a container which is partitioned in plural sections;

b: ~~at least the substances necessary for protein synthesis containing substrate and energy source and a specific translation template are added to the well mentioned in "a"; wherein~~

~~c: the solution in the well mentioned in "b" protein chip reagent is a freeze-dried preparation prepared by freeze drying (by freeze drying);~~

d: an amount of a the deliquescent substance in the freeze-dried preparation ~~in the well mentioned in "c"~~ is less than 0.01 part by weight ~~or less to 1 part~~ by weight of the protein in the said freeze-dried preparation;

~~e: different kind type of translation template is a plurality of different translation templates are contained in each of the different wells of the container the solution mentioned in "b" and makes for making two or more kinds of proteins synthesizable in each different well of the container which is partitioned in plural sections; and~~

~~is~~ a protein synthesized from the translation template is the protein being modified for fixation ~~and is also to a well and/or carrier, the well and/or carrier being~~ coated with a substance having affinity ~~to a substance added by the said modification for fixation to a surface in the well and/or a carrier in the well~~ for the protein that is modified for fixation.

5. (Currently amended): The protein chip reagent utilizing a cell-free protein synthesis system ~~according to~~ of claim 4, wherein the modification for fixation is at least one which is selected from ~~making into avidin, biotin, streptoavidin~~ avidinylation, biotinylation, streptavidinylation, and His tag.

6. (Currently amended): A kit for a cell-free protein synthesis containing the protein chip reagent utilizing a cell-free protein synthesis system ~~mentioned in~~ of claim 1.

7. (Withdrawn): A test method for an interacting substance with a specific protein translated from a specific translation template containing the following elements where a reagent mentioned in claim 1 is used,

(1) a protein chip reagent utilizing a cell-free protein synthesis system is dissolved upon each use;

(2) after dissolving, conditions for a protein translation reaction are regulated to synthesize a specific protein;

(3) a substance to be detected is added and it is confirmed whether the interaction with the specific protein which is synthesized upon each use takes place; and

(4) the interacted substance is judged either qualitatively or quantitatively using a marker.

8. (Currently amended): A kit for a cell-free protein synthesis containing the protein chip reagent utilizing a cell-free protein synthesis system ~~mentioned in~~ of claim 2.

9. (Currently amended): A kit for a cell-free protein synthesis containing the protein chip

reagent utilizing a cell-free protein synthesis system ~~mentioned in~~ of claim 3.

10. (Currently amended): A kit for a cell-free protein synthesis containing the protein chip reagent utilizing a cell-free protein synthesis system ~~mentioned in~~ of claim 4.

11. (Currently amended): A kit for a cell-free protein synthesis containing the protein chip reagent utilizing a cell-free protein synthesis system ~~mentioned in~~ of claim 5.

12. (Withdrawn): A test method for an interacting substance with a specific protein translated from a specific translation template containing the following elements where a reagent mentioned in claim 2 is used,

(1) a protein chip reagent utilizing a cell-free protein synthesis system is dissolved upon each use;

(2) after dissolving, conditions for a protein translation reaction are regulated to synthesize a specific protein;

(3) a substance to be detected is added and it is confirmed whether the interaction with the specific protein which is synthesized upon each use takes place; and

(4) the interacted substance is judged either qualitatively or quantitatively using a marker.

13. (Withdrawn): A test method for an interacting substance with a specific protein translated from a specific translation template containing the following elements where a reagent mentioned in claim 3 is used,

(1) a protein chip reagent utilizing a cell-free protein synthesis system is dissolved upon each use;

(2) after dissolving, conditions for a protein translation reaction are regulated to synthesize a specific protein;

(3) a substance to be detected is added and it is confirmed whether the interaction with

the specific protein which is synthesized upon each use takes place; and

(4) the interacted substance is judged either qualitatively or quantitatively using a marker.

14. (Withdrawn): A test method for an interacting substance with a specific protein translated from a specific translation template containing the following elements where a reagent mentioned in claim 4 is used,

(1) a protein chip reagent utilizing a cell-free protein synthesis system is dissolved upon each use;

(2) after dissolving, conditions for a protein translation reaction are regulated to synthesize a specific protein;

(3) a substance to be detected is added and it is confirmed whether the interaction with the specific protein which is synthesized upon each use takes place; and

(4) the interacted substance is judged either qualitatively or quantitatively using a marker.

15. (Withdrawn): A test method for an interacting substance with a specific protein translated from a specific translation template containing the following elements where a reagent mentioned in claim 5 is used,

(1) a protein chip reagent utilizing a cell-free protein synthesis system is dissolved upon each use;

(2) after dissolving, conditions for a protein translation reaction are regulated to synthesize a specific protein;

(3) a substance to be detected is added and it is confirmed whether the interaction with the specific protein which is synthesized upon each use takes place; and

(4) the interacted substance is judged either qualitatively or quantitatively using a marker.